

Title (en)

GENETIC VARIANTS ON CHR 15Q24 AS MARKERS FOR USE IN DIAGNOSIS, PROGNOSIS AND TREATMENT OF EXFOLIATION SYNDROME AND GLAUCOMA

Title (de)

GENETISCHE VARIANTEN AUF CHR 15Q24 ALS MARKER ZUR VERWENDUNG BEI DER DIAGNOSE, PROGNOSE UND BEHANDLUNG VON EXFOLIATIONSSYNDROM UND GLAUKOM

Title (fr)

VARIANTS GÉNÉTIQUES SUR CHR 15Q24 SERVANT DE MARQUEURS ET DESTINÉS À ÊTRE UTILISÉS DANS LE DIAGNOSTIC, LE PRONOSTIC ET LE TRAITEMENT DU SYNDROME D'EXFOLIATION ET DU GLAUCOME

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Application

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- IS 8668 A 20070808

Abstract (en)

[origin: WO2008152656A2] The present invention relates to methods of diagnosing a susceptibility to an ocular disorder, including glaucoma and exfoliation syndrome. The invention provides methods of diagnosing an increased or decreased susceptibility to exfoliation syndrome and glaucoma, and methods for risk assessment, treatment and prognosis. The invention further relates to kits for use in the methods of the invention.

IPC 8 full level

C12Q 1/68 (2006.01)

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Citation (search report)

See references of WO 2008152656A2

Citation (examination)

- L. A. HINDORFF ET AL: "Genetic architecture of cancer and other complex diseases: lessons learned and future directions", CARCINOGENESIS, vol. 32, no. 7, 31 March 2011 (2011-03-31), pages 945 - 954, XP055045492, ISSN: 0143-3334, DOI: 10.1093/carcin/bgr056
- HAOYU CHEN ET AL: "Ethnicity-based subgroup meta-analysis of the association of LOXL1 polymorphisms with glaucoma", MOLECULAR VISION, vol. 16, 6 February 2010 (2010-02-06), pages 167 - 177, XP055045585

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